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Martin J. King, Ph.D.

Laboratory Director

CLIA ID# 33D1020120

Phys: LABQ MOBILE 42ND ST &amp; LEXINGTON 7616

( ) -  
MARTIN KING PH.D

Patient: ARUN KUMAR, ADITHYAN

DOB: 01/30/2000

Room#

Age:21 Sex:M

Phone: (412) 214-2131

Chart#: 1054134

Route#: 0

Passport: L8869606

Page: 1 of 1

Acc# 2112147549

Coll. Date: 12/28/21

Recv. Date: 12/28/21

Final Report Date: 12/29/21

Order# G-FDC9A124

Coll. Time: 03:54 PM

Recv. Time: 09:00 PM

Final Report Time: 06:25 AM

Test Name

Result

Out of Range

Normal Range

Units

## Report Status: FINAL

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SARS-COV2

NEGATIVE

NEGATIVE

Source: Nasal

This test has been authorized as an EUA for use by authorized laboratories. Negative results do not preclude infection of the upper respiratory tract. Test Methodology: RT-PCR

This SARS-CoV-2 assay is a real time RT-PCR test for the detection of nucleic acid from individuals suspected of having COVID-19 infection. POSITIVE: Results are indicative of active infection with SARS-CoV-2 but do not rule out bacteria infection or co-infection with other viruses.

NEGATIVE: Results do not preclude SARS-CoV-2 infection and should not be used as the sole basis for patient management decisions. If COVID-19 is still suspected based on clinical findings, history and epidemiological information, re-testing of a new specimen should be considered.

Note: Pooling was used to generate this result. Individual specimens with low viral loads may not be detected due to the decreased sensitivity or increased interference when tested with pooled testing.

The United States(U.S.)FDA has made this test available under an Emergency Use Authorization(EUA). The EUA is supported by the Secretary of Health and Human Services (HHS's) declaration that circumstances exist to justify the emergency use of in vitro diagnostics (IVDs) for the detection and/or diagnosis of SARS-CoV-2. This assay has been validated pursuant to the CLIA regulations and may be used for diagnostic testing purposes.

END OF REPORT



Please scan QR Code for verification.

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